Amendments to the Claims:

- 1. (Currently amended) A method for treating pain in a patient in need thereof comprising administering to the patient an amount of substantially enantiomerically pure (S)-norketamine or a pharmaceutically acceptable salt or solvate thereof, which falls in the range of about 0.01 to about 820 mg/kg of body weight of the patient and which, as determined by a physician or medical care provider, is effective to treat pain while not inducing dysphoria.
- 2. (Currently amended) The method of claim 1 in which substantially enantiomerically pure (S)-norketamine is administered to the patient.
 - 3-4. (Canceled)
- 5. (Previously presented)The method of claim 1 in which the amount administered falls in a range of about 1% to about 50% of an amount used to induced anesthesia.
- 6. (Previously presented)The method of claim 1 in which the amount administered falls in a range of about 5% to about 40% of an amount used to induced anesthesia.
- 7. (Previously presented)The method of claim 1 in which the amount administered falls in a range of about 10% to about 20% of an amount used to induced anesthesia.
 - 8. (Canceled)
- 9. (Previously presented) The method of claim 1 in which the amount administered falls in a range of about 0.05 to about 8 mg/kg of body weight of the patient.
- 10. (Currently amended)The method of claim 1 in which a pharmaceutically acceptable salt of substantially enantiomerically pure_(S) noreketamine (S)-norketamine is administered to the patient.
 - 11-12. (Canceled)

- 13. (Previously presented)The method of claim 1 in which the amount is administered over a 24 hour period.
- 14. (Previously presented) The method of claim 1 in which the amount is administered in conjunction with a narcotic analgesic effective to alleviate pain.
- 15. (Currently amended) The method of claim 1, further comprising decreasing a dose of the <u>a</u> narcotic analgesic.
- 16. (Currently amended) A method for self-treating pain in a subject comprising self-administering on an outpatient basis via one or more routes selected from a transmucosal, transdermal, nasal, oral, rectal, vaginal, ocular, or pulmonary route, or any combination of the foregoing routes, an amount of substantially enantiomerically pure (S)-norketamine, or a pharmaceutically acceptable salt or solvate thereof, which falls in the range of about 0.01 to about 820 mg/kg of body weight of the patient and which, as determined by a physician or medical care provider, is effective to treat pain while not inducing dysphoria.
- 17. (Previously presented) The method of claim 16 in which the route of administration is oral.

18-27. (Canceled)

28. (Previously presented) The method of claim 16 in which said pain is selected from the group consisting of breakthrough pain, pain associated with wind-up, chronic pain and neuropathic pain.

29-70. (Canceled)

- 71. (Previously presented)The method of claim 1 in which the amount is administered to the patient via a route selected from the group consisting of intravenous, intramuscular, subcutaneous, intrathecal, and epidural.
 - 72. (Canceled)
- 73. (Withdrawn–currently amended) A method for treating breakthrough pain in a patient in need thereof comprising administering to the patient an amount of substantially

enantiomerically pure (S)-norketamine or a pharmaceutically acceptable salt or solvate thereof, which, as determined by a physician or medical care provider, is effective to treat breakthrough pain while not inducing dysphoria.

- 74. (Withdrawn) The method of claim 73 which further comprises administering a narcotic analgesic.
- 75. (Withdrawn) The method of claim 73 in which the amount is administered orally.
- 76. (Withdrawn) The method of claim 73 in which the amount administered falls in the range of about 0.01 to about 20 mg/kg of body weight of the patient.
- 77. (Withdrawn) The method of claim 73 in which the amount administered falls in the range of about 0.05 to about 8 mg/kg of body weight of the patient.
- 78. (Withdrawn- currently amended) A method for treating pain associated with wind-up in a patient in need thereof comprising administering to the patient an amount of substantially enantiomerically pure (S)-norketamine or a pharmaceutically acceptable salt or solvate thereof, which, as determined by a physician or medical care provider, is effective to treat pain associated with wind-up while not inducing dysphoria.
- 79. (Withdrawn– currently amended) A method for treating chronic pain in a patient in need thereof comprising administering to the patient an amount of substantially enantiomerically pure (S)-norketamine or a pharmaceutically acceptable salt or solvate thereof, which, as determined by a physician or medical care provider, is effective to treat chronic pain up while not inducing dysphoria.
- 80. (Currently amended) A method for treating neuropathic pain in a patient in need thereof comprising administering to the patient an amount of substantially enantiomerically pure (S)-norketamine or a pharmaceutically acceptable salt or solvate thereof, which, as determined by a physician or medical care provider, is effective to treat neuropathic pain up while not inducing dysphoria.

- 81. (Withdrawn– currently amended) An oral dosage form comprising substantially enantiomerically pure (S)-norketamine or a pharmaceutically acceptable salt or solvate thereof and one or more pharmaceutically acceptable excipients, which dosage form, when self-administered in an amount falling in the range of about 0.01 mg/kg to about 20 mg/kg of body weight of the patient, is effective, as determined by a physician or medical care provider, to treat pain while not inducing dysphoria.
- 82. (Withdrawn– currently amended) The oral dosage form of claim 81 which comprises substantially enantiomerically pure (S)-norketamine.
- 83. (Withdrawn– currently amended) The oral dosage form of claim 81 which comprises a pharmaceutically acceptable salt of substantially enantiomerically pure (S)-norketamine.
- 84. (New) A method for treating pain in a patient in need thereof comprising administering orally to the patient an amount of (S)-norketamine or a pharmaceutically acceptable salt or solvate thereof, which falls in the range of about 0.01 to about 8 mg/kg of body weight of the patient and which, as determined by a physician or medical care provider, is effective to treat pain while not inducing dysphoria.
- 85. (New) The method of claim 84 in which the amount administered orally falls in the range of about 0.05 to about 8 mg/kg of body weight of the patient.